

K810251 SYRINGE, ENTMar 11, 1981
41 days to decisionK810251 · Product code: **KYZ** · General HospitalSource: <https://www.510kdatabase.net/k810251/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Irrigating (non Dental) (KYZ)
Date received	Jan 29, 1981
Decision date	Mar 11, 1981
Days to decision	41 days
Third-party review	No

APPLICANT

Company	Megaplast, Inc.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k810251/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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